Lithium lanthanum titanate perovskite ionic conductor: Influence of europium doping on structural and optical properties


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ABSTRACT

The present study reports the insertion of europium ions into the network of lithium lanthanum titanate, La$_{0.55}$Li$_{0.35}$Ti$_O_3$ powders. The materials were prepared efficiently via the Pechini method and were characterized using thermogravimetric analyses, X-ray diffraction, high-resolution electron microscopy and photoluminescence spectroscopy. The results obtained demonstrate the high structural disorder of La$_{0.55}$Li$_{0.35}$Ti$_O_3$ (LLTO) as well as the different environments noticed by Eu$^{3+}$ ions in the La$_{0.55}$Li$_{0.35}$Ti$_O_3$ host lattice. Our results are essentially important once they unfold a better understanding of the complex nature of La$_{0.55}$Li$_{0.35}$Ti$_O_3$ compounds structure.

1. Introduction

La$_{2/3-x}$Li$_{x}$Ti$_O_3$ system has been extensively studied for use in lithium batteries as electrodes, solid electrolytes and as separators in lithium-air batteries as a result of its high ion mobility, which can reach up to $10^{-3}$ S cm$^{-1}$ at room temperature [1–3]. Based on the work conducted by Inaguma et al. [4] which reported high mobility of lithium ions in La$_{0.75}$Li$_{0.25}$Ti$_O_3$ composition, several mechanisms for the conduction of La$_{2/3-x}$Li$_{x}$Ti$_O_3$ compounds have since been proposed associated mainly with structural considerations, conductivity measurements and theoretical models aiming at justifying the high ionic conductivity at room temperature [5–14]. Despite considerable understanding has been achieved in LLTO compounds, detailed comprehension of lithium ion diffusion as well as the influence of composition and associated local structural changes on diffusion is still lacking [14]. In fact since the different crystal structures obtained (including simple cubic [15–17], hexagonal [18], tetragonal [19–21] and orthorhombic [21,22]) depend on the method of synthesis used, and the conductivity is related to the structure, it is possible to control the conductivity indirectly via the synthesis.

Most of the authors assume that the high ionic conductivity stems from the presence of vacancies on La$_{2/3-x}$Li$_{x}$Ti$_O_3$ sites, favoring the mobility of Li$^+$ ions through the structure. Thus, the position of Li$^+$ ions is of crucial importance in explaining the ionic conductivity in La$_{2/3-x}$Li$_{x}$Ti$_O_3$ compounds which is directly related to the occupancy of the interstitial spaces by Li$^+$ [23].

The replacement of the elements A or B from the ABO$_3$ perovskite structure of La$_{2/3-x}$Li$_{x}$Ti$_O_3$ is carried out aiming at improving the ionic conductivity of these materials. A number of studies have reported a complete or partial substitution of La$^{3+}$ with other lanthanide ions including Pr, Nd and Sm, where it was observed that this substitution is accompanied by distortions in the perovskite lattice. These distortions are known to be responsible for the reduction of the so-called "bottlenecks". As the bottlenecks are the path by which Li ions move, their reduction is seen to account for the decrease in ionic conductivity [23].

Similarly, the lanthanide Eu$^{3+}$ is found to be capable of replacing La$^{3+}$ ions from La$_{2/3-x}$Li$_{x}$Ti$_O_3$ and be used as structural probe. The electronic transitions located in their luminescent centers indicate the ionic properties of the matrix in which it is contained as well as the interaction Eu$^{3+}$/matrix. Hence, the use of lanthanides as structural probe provides information regarding the symmetry of the crystallographic sites, the nature of the chemical bonds (covalent grade), the energy transfer processes and the long range symmetry effect of the spectroscopic properties of the luminescent centers [24–31].

In this work, the replacement of La$^{3+}$ with Eu$^{3+}$ was done with the purpose of understanding the complex structure of La$_{0.55}$Li$_{0.35}$Ti$_O_3$ and the location of the ions in the crystal lattice for a better knowledge of the ionic conductivity in La$_{2/3-x}$Li$_{x}$Ti$_O_3$.

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2. Methodology

La0.5Li0.5TiO3 powders which are nominally pure and Eu3+-doped (0.1, 0.25, 0.5 and 1 at%) were obtained by the Pechini method. Titanium (IV) isopropoxide, Ti(OCH2CH3)4 (Alfa Aesar), lithium nitrate, LiNO3 (Vetec), lanthanum nitrate, La(NO3)3·6H2O (Vetec), europium oxide, Eu2O3 (Sigma Aldrich), citric acid, C6H8O7 (Sigma Aldrich) and ethylene glycol, C2H6O2 (Sigma Aldrich) were used for the synthesis. The molar ratio of the chemicals was defined based on the Pechini method [32]: 1 mol of metal: 4 mol of citric acid: 16 mol of ethylene glycol. The precursors were mixed, stirred and heated at 120 °C for 1 h, followed by heat treatment at 350 °C for 2 h (precursor powders). The final material was slowly burned at 800 °C for 3 h. The precursor powders were characterized by thermogravimetry (TG) and differential thermal analysis (DTA) using a Netsch-Thermische Analyse (PU 1.851.01 and TASC 414/2 controller). α-Al2O3 was used as a standard material and the samples were placed in alumina crucibles using a 10 Pt thermocouple (Pt / Pt 10% Rh) and subjected to a heating rate of 10 °C/min and synthetic air flow of 30 cm3/min. The X-ray diffraction (XRD) measurements were performed using a Rigaku/RINT2000, 2θ = 30.2° and TASC 414/2 controller). The FEG-VP Zeiss Supra 35 model was used for the FE-SEM (high resolution field emission gun scanning electron microscopy) measurements. Optical measurements were performed by photoluminescence (PL) using a Thermal Jarrell-Ash Monospec 27 monochromator and a Hamamatsu R446 photomultiplier. A krypton ion laser (Coherent Innova) with an exciting wavelength of 350 nm was used, with the nominal output power of the laser kept at 550 mW. UV-Vis measurements were performed using a LAMBA 1050 PerkinElmer spectrophotometer and Raman spectra were collected with a triple monochromator T-64000 HORIBA Jobin-Yvon (USA) coupled to a CCD detector with excitation of 514.4 nm from an argon ion laser and input power of 8 mW. All measurements were performed at room temperature.

3. Results and discussion

The heating process of the polymeric precursor method involves a great loss of mass. TG curve of La0.5Li0.5TiO3 precursor powders (Fig. 1a) indicates a mass loss of approximately 70% between 25 °C and 900 °C, while the curve of La0.5Li0.5TiO3: Eu3+ (Fig. 2a) shows a mass loss of approximately 60% between 25 °C and 900 °C. The mass loss during the heat treatment refers to the removal of solvent and water, the decomposition of the polyester chain and carboxyl groups linked to the metal with a consequent formation of crystalline metal-oxygen phase.

DTA curves (Fig. 1a and Fig. 2a) show both endothermic and exothermic processes involved in the decomposition of the polyester. Endothermic processes, for temperatures below 200 °C, are associated with the loss of the excess of ethylene glycol and water. The exothermic process is seen beginning at 450 °C. A maximum heat is found to be released at 500 °C relating to the decomposition of the polyester chain. The formation of the metal-oxygen bonds is observed from 500 °C onwards, giving rise to the formation of the perovskite phase of lithium lanthanum titanate - L2O.5Li0.5TiO3.

Figs. 1b and 2b present FE-SEM images of the non-doped and doped materials, respectively. These images reveal aggregated nanoparticles with a spherical shape. It can be observed that the doped powder has larger particles, which are found to be more agglomerated and with sintering traces compared to the non-doped powders. The addition of Eu3+, in effect, promotes distortions and instability in the network frame, thus favoring the sintering process.

The XRD patterns of non-doped La0.5Li0.5TiO3 powders are shown in Fig. 1c. The powders treated at 350 °C are amorphous. The formation of La0.5Li0.5TiO3 crystalline phase (JCPDS 01-070-6720) can be observed at 700 °C. The powders heated at 800 °C are found to exhibit high crystallinity and this outcome suggests that in the temperature range studied, the increase observed in the annealing temperature improves the crystallinity of the perovskite phase. It is therefore evidently clear the existence of a correlation between the crystallization processes of La0.5Li0.5TiO3 phase and the removal of the organic components. A careful analysis of the diffraction patterns revealed the presence of the secondary phase of Li2TiO3 (JCPDS 00-033-0831) denoted in the diffraction patterns with the symbol (*) at 2θ = 30.2° and 2θ = 43.5°.

The XRD results of the Eu3+ doped powders are shown in Fig. 2c. It is worth pointing out that no changes were observed in the diffractions peaks when La0.5Li0.5TiO3 was doped with Eu3+ as compared to the nominally pure material; Eu3+ was inserted in the crystalline network of the La0.5Li0.5TiO3 and no secondary phase regarding Eu3+ doping was observed. Furthermore, no changes are observed in the XRD of the material doped with varying concentrations of Eu3+ ion.

Raman measurements were evaluated in order to obtain additional information regarding the short range structural ordering of LLTO samples. The spectra (Fig. 3) show seven bands located at 140, 230, 360, 400, 530 and 660 cm−1. The bands around 140, 230 and 360 cm−1 can be assigned to Eg modes while the A1g mode is found at 230 cm−1. According to Sanjuán et al. [34] the bands near to 140 and 320 cm−1 are present in the spectra mostly in plane and c-axis titanium vibration, respectively, and the other modes exist due to the oxygen motion. Bands located around 360, 400 and 660 cm−1 are present in lithium titanates Raman spectra [35,36] thus the presence of these bands in our Raman spectra is in agreement whit our XRD results, which revealed the presence of Li2TiO3 as a secondary phase. Regarding
the Eu-doped samples, it was possible to observe a decrease of intensity compared to the non-doped samples, which is more evident in the bands located at 360 and 400 cm\(^{-1}\) attributed to the lithium titanate, that almost disappear. Besides that, the presence of europium also promotes the displacement of the bands. This behavior is due to the substitution of A-site by Eu\(^{3+}\) which could affect the motion in the TiO\(_6\) octahedra, as observed to FeO\(_6\) octahedra in rare earth doped BiFeO\(_3\) perovskite [37]. According to Ye et al. in the study of a double-perovskite, the A-site substitution by Eu\(^{3+}\) gives a shift of the peaks, even though the substituted content was small (x ≤ 0.10). When the B-sites are substituted no clear peak shift is observed [38].

The optical band gap energy of the samples was calculated by the application of Wood-Tauc method [38] and the Kubelka-Munk function [39]. This methodology is used to transforms diffuse reflectance measurements into E gap values with high accuracy [40]. Fig. 4, shows the E gap values of pure LLTO and Eu\(^{3+}\)-doped LLTO, where it is possible to note a slight decrease in the optical band gap with the incorporation of Eu\(^{3+}\). The same behavior could be noticed for non-doped ZnO and Eu\(^{3+}\)-doped ZnO [41]. The different E gap values obtained indicates the existence of intermediary energy levels between the valence band and conduction band. The increase of the optical band gap values is related to a reduction of the intermediary energy levels between bands due to the reducing the number of defects in the structure of the material, since the optical band gap energy is controlled by the degree of structural order-disorder in the lattice [40]. In this sense, the increase of disorder structural is related to the Eu\(^{3+}\) content, since we can suppose that the more ordered sample is pure LLTO due to the higher band gap value, suggesting a different conformation or a low concentration of energy states between bands [40]. These energy levels are related to structural order-disorder which causes the loss of symmetry between O–M–O bonds (M = metal) resulting in complex clusters formation [40].

Fig. 5a shows the emission spectrum of non-doped La\(_{0.5}\)Li\(_{0.5}\)TiO\(_3\). PL analyses provide important information concerning the electronic structure of the materials. Here, we can observe a broadband covering the visible electromagnetic spectra within the range of 400–600 nm centered at 450 nm (blue emission). This feature is likely to be associated with the presence of defects known to be typical of multiphonon and multilevel emission processes [42]. The broadband model suggests that the PL emission presented by semiconductor materials results of electronic transitions due to the distortion in the crystalline lattice [40]. Non-doped La\(_{0.5}\)Li\(_{0.5}\)TiO\(_3\) presents structural disorder in the material's lattice promoting the formation of energy states within the band gap, contributing to the broadband emission in their photoluminescence spectra.

PL emission may be attributed to distortions on the lattice, cation or anion vacancies, surface states and quantum confinement [43]. Emission at green/yellow/orange and red region is caused by deep holes and shallow holes are responsible for PL emission at violet and blue region [44,45]. Deep levels are generated by intermediary states close to the conduction band, while shallow levels are due to states near the valence band [43].

Thus, in the case of non-doped La\(_{0.5}\)Li\(_{0.5}\)TiO\(_3\), the PL band centered at 450 nm may essentially indicate the contribution of defects results from shallow levels to PL emission. This information may be backed by the deconvolution of the PL band (Fig. 5b) which was carried out using PeakFit program [31] and the Voigt function. It is possible to observe that the major PL band is composed of three bands, two of them corresponding to different energies of violet-blue emission (440 nm and 478 nm), while the other one is related to emission in green region (515 nm). Interestingly, this is additional evidence that the shallow defects are responsible for the emission observed in the non-doped La\(_{0.5}\)Li\(_{0.5}\)TiO\(_3\) spectrum, since the violet-blue region occupied 80% of the major PL band coupled with the fact that there is low contribution of emission characteristic of deep defects (20%). Since PL emission in semiconductors is results of electronic transition caused by distortion in the crystalline lattice, this behavior can be associated with the lattice distortion caused by the difference in size of Li and La atoms which
generate shallow states close to the valence band.

The analysis of the emission spectra of Eu$^{3+}$ doped materials (Fig. 6a) reveals that the emissions related to the matrix are observed only at lower Eu$^{3+}$ concentrations. The increase in the Eu$^{3+}$ content leads to the disappearance of the emission band attributed to the La$_{0.5}$Li$_{0.5}$TiO$_3$ matrix. When the Eu$^{3+}$ is present in a greater amount, the relaxation via excitation occurs preferably through the dopant once the Eu$^{3+}$ emission is favored instead of the matrix emission.

Eu$^{3+}$ doped compounds present intraconfigurational emissions associated with $4\ell^6 \rightarrow 4\ell^6$ transitions (Fig. 6b). Regarding the excited level $^5D_0$, the lines observed in the spectrum are known to be $^5D_0 \rightarrow ^7F_J$ ($J = 0, 1, 2, 3, 4$). The lines with $J = 5$ and 6 are found to have low intensity and as such were not observed. A further observation worth mentioning is the emission from the higher excited state $^5D_1 \rightarrow ^7F_J$ ($J = 0, 1, 2$). However, transitions starting from $^5D_2$ and $^5D_3$ excited levels were not observed; when Eu$^{3+}$ ions are excited to any level above $^5D_1$, a fast non-radiative multiphonon relaxation to these levels occurs. Table 1 lists the Eu$^{3+}$ ions transitions in a La$_{0.5}$Li$_{0.5}$TiO$_3$ host lattice and the respective wavelength of the emission band.

Given the fact that the levels involved in the transitions are derived
from the same configuration (4f6), it would be quite expected that all these transitions were forbidden. Nevertheless, the ligand field (odd) associated with the chemical environment where the Eu3+ ion is located has the property of mixing levels of opposite parity settings, relaxing the selection rules for the electric dipole transitions. This fact is reflected in all 5D0 → 7F(J) (0-J) transitions, being more noticeable in the transition 5D0 → 7F2 (0−2), which is known to be hypersensitive; and excluding 5D0 → 7F1 (0−1) which is a magnetic transition [24–30].

The intensity of 5D0 → 7F1 (0-J) transitions essentially depends on the symmetry around the Eu3+ ions. When Eu3+ is inserted in high symmetry environments, magnetic dipole transitions 5D0 → 7F1 (0−1) are found to be predominant. Contrary to that, if Eu3+ occupies a non-symmetric environment (in other words of low symmetry), electric dipole transitions become largely predominant [30]. As shown in Fig. 4a, the (0−2) transitions are relatively more intense (0−1) for all the samples, indicating that Eu3+ doping exhibits a characteristic profile of dopants occupying non-symmetric centers.

The intensity ratio (0−2)/(0−1) is denoted as asymmetry ratio R. The higher the value of R, the larger will be the magnitude of the electric dipole transition and low ligand symmetry, leading to stronger...
splitting of transitions. The smaller the R value, the more symmetric is the Eu\(^{3+}\) environment [29]. The asymmetry ratio R can likewise be used to investigate the relative strength of covalent or ionic bonding of the Eu\(^{3+}\) ions and the surrounding ligands. The higher the value of R, the higher the presence of covalent bond and vice versa [29]. In our case, the average of the calculated R for all the doped samples was ~1.65, confirming the distortion of the local symmetry. Increasing the concentrations of Eu\(^{3+}\) from 0.1 to 0.5 at% leads to a slight increase in R, from ~1.64 to ~1.74, respectively. However, the sample doped with 1 at% Eu\(^{3+}\) presented the lowest R, 1.52. Fig. 7 illustrates R as a function of Eu\(^{3+}\) content.

It is known from the complex structure of La\(_{0.5}\)Li\(_{0.5}\)TiO\(_3\) that there are many different types of environments that Eu\(^{3+}\) could occupy. The results from photoluminescence spectroscopy suggest that Eu\(^{3+}\) when inserted in the La\(_{0.5}\)Li\(_{0.5}\)TiO\(_3\) matrix occupies at least three different chemical environments. Fig. 8 illustrates the different possible settings that Eu\(^{3+}\) may be occupying.

La\(_{0.5}\)Li\(_{0.5}\)TiO\(_3\) is an ABO\(_3\) type perovskite where the substitution of divalent ions located in sites A with trivalent ions such as La\(^{3+}\) gives rise to vacancies on sites A. When a percentage number of La\(^{3+}\) ions was replaced by Li\(^+\) ions, we expected to see the Li\(^+\) ions occupying sites A of the structure, with a coordination of 12. Admittedly, Li\(^+\) is too small to favorably occupy these sites, instead, Li\(^+\) appears to be occupying the interstices of the structure, which are capable of accommodating a large particle size aside from being more agglomerated when doped, suggesting that the addition of Eu\(^{3+}\) favors the sintering process of the LLTO matrix as a dopant.

By means of morphological analysis, the powder was found to have a large particle size aside from being more agglomerated when doped, suggesting that the addition of Eu\(^{3+}\) favors the sintering process of La\(_{0.5}\)Li\(_{0.5}\)TiO\(_3\), which was explained by the introduction of defects into the lattice of the material. This result is in accordance with the UV–Vis measurements, which revealed a small decrease in the gap of the doped materials.

The ratio between the transitions (0→2) and (0→1) obtained by photoluminescence spectroscopy revealed a characteristic profile of Eu\(^{3+}\) occupying low symmetry sites. Furthermore, no changes were observed in the site occupied by Eu\(^{3+}\) in spite of a change in the concentration of the dopant. Our results show that La\(^{3+}\) in the La\(_{0.5}\)Li\(_{0.5}\)TiO\(_3\) structure is largely influenced by different types of atoms and neighboring defects.

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